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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Group: 1626

Confirmation No.: 9842

Application No.: 10/725,191

Invention: **PHOSPHORAMIDE
COMPOUNDS**

Applicant: Richard F. Borch et al.

Filed: December 1, 2003

Attorney Docket: 3220-200986

Examiner: Golam M. Shameem

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} I hereby certify that this correspondence is being
} filed electronically in the United States Patent and
} Trademark Office

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}
} on March 23, 2007

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} (Signature)

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}
} Charles L. Twyman
} (Printed Name)

SUBMISSION MADE WITH AN RCE:

Supplemental Information Disclosure Statement

Mail Stop RCE
Commissioner for Patents
P. O. Box 1450
Alexandria, Virginia 22313-1450

Sir:

Applicants have concurrently filed a request for continued examination (RCE) under 37 C.F.R. § 1.114 in a separate paper. In association with that RCE, Applicants hereby request entry and consideration of the instant Submission, which Submission includes a Supplemental Information Disclosure Statement (IDS), Form 1449, and all required copies of the references cited therein.

This IDS is filed in the application identified above pursuant to 37 C.F.R. § 1.56. No representation is intended that a complete search has been made of the prior art or that no better art references than the references cited in the IDS are available. The filing of this IDS shall not be construed to be an admission that the information cited in the IDS is, or

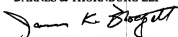
1.98(a)(2)(ii), copies of the cited U.S. patents are not provided herewith. Copies of the cited foreign patent documents and other references (non-patent) have been provided.

The cited references are not believed to disclose or suggest the invention recited in the claims of the above-identified application. It is therefore believed that the claimed invention is patentably distinguishable over the cited references.

Applicants have authorized, concurrent to filing the instant IDS, Barnes & Thornburg LLP Deposit Account 10-0435 to be debited in the amount of \$575.00, which amount is the sum of the \$180.00 fee set forth under 37 C.F.R. 1.17(p) for filing an Information Disclosure Statement, and the \$395.00 fee set forth under 37 C.F.R. § 1.17(e) for requesting continued examination under small-entity status.

Applicants do not believe that any additional fees are due for filing the instant Information Disclosure Statement. However, the Commissioner is hereby authorized to charge any additional fees that may be required, or credit any overpayment, to Barnes & Thornburg LLP Deposit Account No. 10-0435, with reference to our matter number 3220-200986.

Respectfully submitted,
BARNES & THORNBURG LLP



James K. Blodgett
Registration No. 48,480
Agent for Applicants

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317-231-7401

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE INFORMATION DISCLOSURE STATEMENT				ATTY. DOCKET NO.: 3220-200986		SERIAL NO.: 10/725,191	
				APPLICANT(S): Borch et al.			
				FILING DATE: Dec. 1, 2003		GROUP: 1626	

U.S. PATENT DOCUMENTS							
*Examiner Initial	Document Number	Date	Name	Class	Subclass	Filing Date if appropriate	
	AA	5,233,031					
	AB	6,903,081					
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	AK						

FOREIGN PATENT DOCUMENTS							
	Document Number	Date	Country	Class	Subclass	Translation Yes No	
	AL	WO 93/06120	April 1, 1993	PCT			
	AM	WO 01/74827	Oct. 11, 2001	PCT			
	AN						
	AO						

OTHER REFERENCES (Including Author, Title, Date, Pertinent Pages, Etc.)	
AR	"The Synthesis of O-Monosaccharidyl-Methoxycarbonyl-Phosphoramidates by Arbuzov Reaction," Chen et al., Chinese Chemical Letters, vol. 6, No. 1, pp. 23-26, 1995
AS	"Design and Synthesis of Lipophilic Phosphoramidate d4T-MP Prodrugs Expressing High Potency Against HIV In Cell Culture: Structural Determinants for <i>in vitro</i> Activity and QSAR," Steelquist et al., J. Med. Chem. 1998, 42, 4122-4128
AT	"Phosphoramidates as Potent Prodrugs of Anti-HIV Nucleotides: Studies in the Amino Region," McGuigan et al., Antiviral Chemistry & Chemotherapy (1996) 7(1), 31-36
AU	"Design and Synthesis of Novel Nucleotide Prodrugs," Meyers et al., American Association for Cancer Research 2000 (Abstract Proof Page), Abstract #100710, dated December 2, 1999
AV	"Protein Tyrosine Kinases and Cancer," Kolibaba et al., "Biochimica et Biophysica Acta 1333 (1997) F217-F248
AW	"Receptor Protein-Tyrosine Kinases and Their Signal Transduction Pathways," van der Geer et al., Annu. Rev. Cell Biol., 1994, 10:251-337
AX	"Src Homology Region 2 Domains Direct Protein-Protein Interactions in Signal Transduction," Moran et al., Proc. Natl. Acad. Sci. USA, Vol. 87, Nov. 1990, pp. 8622-8626
AY	"Binding of SH2 Domains of Phospholipase C γ 1, GAP, and Src to Activated Growth Factor Receptors," Anderson et al., Science, Vol. 250, Nov. 16, 1990, 979-982
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Examiner	Date Considered
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*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 809.
 Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE				ATTY. DOCKET NO.: 3220-200986		SERIAL NO.: 10/725,191	
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U.S. PATENT DOCUMENTS							
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FOREIGN PATENT DOCUMENTS							
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BR	"Recognition and Specificity in Protein Tyrosine Kinase-Mediated Signalling," Songyang et al., TIBS 20, November 1995, 470-475
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BT	"Peptide Inhibitors of src SH3-SH2-Phosphoprotein Interactions," Gilmer et al., The Journal of Biological Chemistry, Vol. 269, No. 50, December 16, 1994, 31711-31719
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BV	"Design, Synthesis, and Cocrystal Structure of a Nonpeptide SrcSH2 Domain Ligand," Plummer et al., J. Med. Chem. 1997, 40, 3719-3725
BW	"Inhibition of SH2 Domain/Phosphoprotein Association by a Nonhydrolyzable Phosphonopeptide," Domchek et al., Biochemistry 1992, 31, 9865-9870
BX	"Nonhydrolyzable Phosphotyrosyl Mimetics for the Preparation of Phosphatase-resistant SH2 Domain Inhibitors," Burke, Jr., et al., Biochemistry, 1994, 33, 6490-6494
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BZ	"Conformationally Constrained Phosphotyrosyl Mimetics Designed as Monomeric Src Homology 2 Domain Inhibitors," Burke Jr., et al., J. Med. Chem. 1995, 38, 1386-1396

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 Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

BASED ON FORM PTO 1449

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE				ATTY. DOCKET NO.: 3220-200986		SERIAL NO.: 10/725,191	
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U.S. PATENT DOCUMENTS				GROUP: 1626			
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	CA						
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FOREIGN PATENT DOCUMENTS							
		Document Number	Date	Country	Class	Subclass	Translation Yes No
	CL						
	CM						
	CN						
	CO						
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	CR	*4'-O-[2-(2-Fluoromethyl)-L-Tyrosine: A Phosphotyrosyl Mimic for the Preparation of Signal Transduction Inhibitory Peptides," Burke Jr., et al., J. Med. Chem. 1996, 39, 1021-1027					
	CS	*Phosphotyrosine-Containing Dipeptides as High-Affinity Ligands for the p56 ^{lck} SH2 Domain," Llinas-Brunat et al., J. Med. Chem. 1999, 42, 722-729					
	CT	*Acquisition of High-Affinity, SH2-Targeted Ligands via a Spatially Focused Library," Lee et al., J. Med. Chem. 1999, 42, 784-787					
	CU	*Ligands for the Tyrosine Kinase p56 ^{lck} SH2 Domain: Discovery of Potent Dipeptide Derivatives with Monocharged, Nonhydrolyzable Phosphate Replacements," Beaulieu et al., J. Med. Chem. 1999, 42, 1757-1766					
	CV	*Synthesis and Biological Evaluation of 5-Fluoro-2'-Deoxyuridine Phosphoramidate Analogs," Fries et al., J. Med. Chem., 1995, 38, 2672-2680					
	CW	*Synthesis and Biological Activity of Novel 5-Fluoro-2'-Deoxyuridine Phosphoramidate Prodrugs," Meyers et al., J. Med. Chem., 2000, 43, 4313-4318					
	CX	*Synthesis and Evaluation of Nitroheterocyclic Phosphoramidates as Hypoxia-Selective Alkylating Agents," Borch et al., J. Med. Chem., 2000, 43, 2258-2265					
	CY	Gibbs et al., Current Medicinal Chemistry, Vol. 8, No. 12, October 2001, pp. 1437-1465					
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	DA						
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FOREIGN PATENT DOCUMENTS							
		Document Number	Date	Country	Class	Subclass	Translation Yes No
	DL						
	DM						
	DN						
	DO						

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	DR	Weldong et al., American Chemical Society. Abstracts of Papers at the National Meeting, ACS, Washington, D.C., Vol. 228, Part 1, 26 August 2004, p. U930
	DS	
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